

# Salmoporc, lyophilisate for oral suspension for pigs

Authorised

- *Salmonella enterica*, subsp. *enterica*, serovar *Typhimurium*, strain 421/125 (histidine-adenine auxotrophic), Live

## Product identification

**Medicine name:**

Salmoporc lyophilisate for oral suspension for pigs  
Salmoporc, lyophilisate for oral suspension for pigs

**Active substance:**

*Salmonella enterica*, subsp. *enterica*, serovar *Typhimurium*, strain 421/125 (histidine-adenine auxotrophic), Live

**Target species:**

Pig

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

*Salmonella enterica*, subsp. *enterica*, serovar *Typhimurium*, strain 421/125 (histidine-adenine auxotrophic), Live

8.00 log10 colony forming unit(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Oral lyophilisate

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**Withdrawal period by route of administration:**

**Oral use:**

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**Pig**

- Meat and offal. 6 week

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI09AE02

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Ireland

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**Package description:**

Cardboard box containing one 10 ml glass vial (type I) with rubber stoppers and aluminium crimp caps containing 200 doses (1 dose = 1 ml) of lyophilisate.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Similar biological application (Article 13(4) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ceva Sante Animale

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**Marketing authorisation date:**

18/04/2019

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**Manufacturing sites for batch release:**

Ceva-Phylaxia Veterinary Biologicals Co. Ltd  
IDT Biologika GmbH

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA10815/064/002

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**Date of authorisation status change:**

18/04/2019

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0247/002

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**Concerned member states:**

Austria Belgium Czechia Denmark Hungary Ireland Italy Portugal Romania  
Slovakia United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to

[www.adrreports.eu/vet](http://www.adrreports.eu/vet)